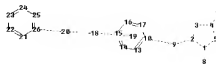
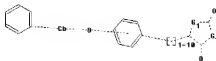


=>

Uploading C:\Program Files\Stnexp\Queries\10530757-genus-F.str



chain nodes :

7 8 9 18 20

ring nodes :

1 2 3 4 5 10 13 14 15 16 17 21 22 23 24 25 26

chain bonds :

1-8 2-9 4-7 9-10 18-20 20-26

ring bonds :

1-2 1-5 2-3 3-4 4-5 10-13 10-17 13-14 14-15 15-16 16-17 21-22 21-26

22-23 23-24 24-25 25-26

exact/norm bonds :

1-2 1-5 1-8 2-3 2-9 3-4 4-5 4-7 9-10 18-20 20-26

normalized bonds :

10-13 10-17 13-14 14-15 15-16 16-17 21-22 21-26 22-23 23-24 24-25 25-26

isolated ring systems :

containing 1 :

G1:O,S,N

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 7:CLASS 8:CLASS 9:CLASS 10:Atom 13:Atom

14:Atom 15:Atom 16:Atom 17:Atom 18:CLASS 19:Atom 20:Atom 21:Atom 22:Atom

23:Atom 24:Atom

25:Atom 26:Atom

Generic attributes :

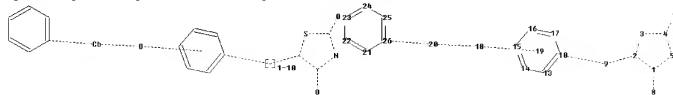
20:

Saturation : Unsaturated

L1 STRUCTURE UPLOADED

=>

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chain nodes :

```

7  8  9 18 20
ring nodes :
1  2  3  4  5 10 13 14 15 16 17 21 22 23 24 25 26
chain bonds :
1-8 2-9 4-7 9-10 18-20 20-26
ring bonds :
1-2 1-5 2-3 3-4 4-5 10-13 10-17 13-14 14-15 15-16 16-17 21-22 21-26
22-23 23-24 24-25 25-26
exact/norm bonds :
1-2 1-5 1-8 2-3 2-9 3-4 4-5 4-7 9-10 18-20 20-26
normalized bonds :
10-13 10-17 13-14 14-15 15-16 16-17 21-22 21-26 22-23 23-24 24-25 25-26

```

```

isolated ring systems :
containing 1 :

```

G1:O,S,N

```

Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 7:CLASS 8:CLASS 9:CLASS 10:Atom 13:Atom
14:Atom 15:Atom 16:Atom 17:Atom 18:CLASS 19:Atom 20:Atom 21:Atom 22:Atom
23:Atom 24:Atom
25:Atom 26:Atom
Generic attributes :
20:
Saturation           : Unsaturated

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L2 STRUCTURE UPLOADED

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FILE 'REGISTRY' ENTERED AT 13:00:15 ON 26 JUN 2008
L1 STRUCTURE UPLOADED
L4 5 S L1 SSS FULL

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FILE 'CAPLUS' ENTERED AT 13:01:19 ON 26 JUN 2008
L5 1 S L4

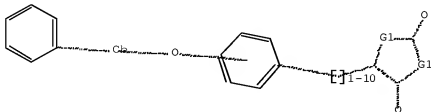
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FILE 'REGISTRY' ENTERED AT 13:01:26 ON 26 JUN 2008

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=> d l1
L1 HAS NO ANSWERS
L1 STR

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G1 O, S, N

Structure attributes must be viewed using STN Express query preparation.

=> fil caplus

=> d 15 bib abs

L5 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2008 ACS on STN

AN 2003:757334 CAPLUS Full-text

DN 139:276885

TI Preparation of novel heterocyclic analogs of diphenylethylene compounds as antidiabetics

IN Neogi, Partha; Dey, Debendranath; Medicherla, Satyanarayana; Nag, Bishwajit; Lee, Arthur

PA USA

SO U.S. Pat. Appl. Publ., 66 pp., Cont.-in-part of U.S. Ser. No. 843,167. CODEN: USXXCO

DT Patent

LA English

FAN.CNT 14

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 20030181494	A1	20030925	US 2002-265902	20021008
	US 20020025975	A1	20020228	US 2001-785554	20010220
	US 20020032225	A1	20020314	US 2001-843167	20010427
	US 7105552	B2	20060912		
	CA 2501456	A1	20040422	CA 2003-2501456	20031008
	WO 2004033438	A1	20040422	WO 2003-US31803	20031008
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	RW:				
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	AU 2003282754	A1	20040504	AU 2003-282754	20031008
	EP 1549625	A1	20050706	EP 2003-774638	20031008
	R:				
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	CN 1708486	A	20051214	CN 2003-80101164	20031008
	JP 2006050551	T	20060216	JP 2004-543490	20031008
	NZ 539121	A	20080430	NZ 2003-539121	20031008
	US 20060235062	A1	20061019	US 2006-530757	20060508
PRAI	US 1999-287237	A2	19990406		
	US 2000-591105	B2	20000609		
	US 2001-785554	A2	20010220		
	US 2001-843167	A2	20010427		
	US 1998-74925	A2	19980508		
	US 2002-265902	A	20021008		
	WO 2003-US31803	W	20031008		
OS	MARPAT 139:276885				
GI					

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB The title compds. [I; Z = II-IV; n, m, q and r = 0-4 ($n+m \leq 4$ and $q+r \leq 4$); p, s = 0-5 ($p+s \leq 5$); R, R2 = H, alkyl, alkenyl, etc.; R1 = H, alkyl, alkenyl, etc.; A, A1, A2 = H, acylamino, acyloxy, alkanoyl, etc.; B, B1, B2 = H, acylamino, acyloxy, alkanoyl, etc.; or A and B together, or A1 and B1 together, or A2 and B2 together, may be joined to form a methylenedioxy or ethylenedioxy; X, X1 = (un)substituted NH, O, S] which are effective in lowering blood glucose level, serum insulin, triglyceride and free fatty acid levels in animal models of Type II diabetes, were prepared E.g., a multi-step synthesis of V, starting from 3,5-dimethoxybenzaldehyde and 4-hydroxyphenylacetic acid, was given. The compound V showed strong glucose lowering activity even though it is a weak PPAR- γ agonist (data given). The compds. I are disclosed as useful for a variety of treatments including the treatment of inflammation, inflammatory and immunol. diseases, insulin resistance, hyperlipidemia, coronary artery disease, cancer and multiple sclerosis. Pharmaceutical composition comprising the compound I was claimed.

SESSION WILL BE HELD FOR 120 MINUTES

STN INTERNATIONAL SESSION SUSPENDED AT 13:01:56 ON 26 JUN 2008